

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) EP 1 061 555 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
20.12.2000 Bulletin 2000/51

(51) Int Cl.7: H01J 63/06, H01J 9/24

(21) Application number: 00305140.6

(22) Date of filing: 16.06.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 18.06.1999 KR 9923051
02.06.2000 KR 0030354

(71) Applicants:
• Iljin Nanotech Co., Ltd.
Seoul (KR)
• Lee, Cheol-jin
Gunsan-city, Jeonlabuk-do (KR)
• Cho, Young-sang
Incheon (KR)

(72) Inventors:
• Lee, Cheol-jin
Gunsan-city, Jeonlabuk-do (KR)
• Cho, Young-sang, 602-1008 New Seoul Apt.
Incheon (KR)
• Yoo, Jae-eun, 106-1001 Kyungnam Apt.
Seoul (KR)

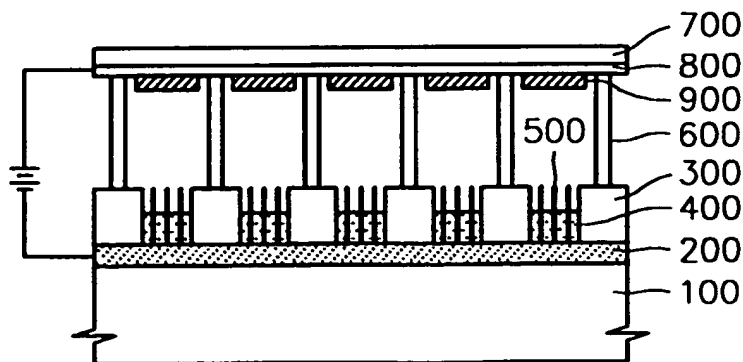
(74) Representative: Stuttard, Garry Phillip
Urquhart-Dykes & Lord
Tower House
Merrion Way
Leeds LS2 8PA (GB)

(54) White light source using carbon nanotubes and fabrication method thereof

(57) A white light source using carbon nanotubes and a method of fabricating the same are provided. The white light source includes a metal film (200) which is used as a cathode and formed on a lower substrate (100), a conductive polymer film pattern (400) formed on the metal film, carbon nanotubes (500) which are substantially vertically bound with the conductive poly-

mer film pattern such that one end thereof is exposed above the surface of the conductive polymer film pattern, for emitting electrons, spacers (600) installed on the metal film, and a transparent upper substrate (700) which has a transparent electrode (800), to which a fluorescent body (900) is attached, and is mounted on the spacers such that the fluorescent body faces the carbon nanotubes.

FIG. 1



EP 1 061 555 A1

Description

BACKGROUND OF THE INVENTION

1. Field of the Invention

[0001] The present invention relates to a white light source, and more particularly, to a method of manufacturing a white light source having an excellent luminous efficacy.

2. Description of the Related Art

[0002] A representative white light source is a fluorescent lamp. The fluorescent lamp uses the emission of light by a fluorescent body due to a discharge effect. This fluorescent lamp has the drawback of low luminance. Moreover, it is difficult to miniaturize the fluorescent lamp and to make the fluorescent lamp operate with low operating voltage. In addition, as use time lapses, the luminance of the fluorescent lamp decreases. Consequently, the stability and reliability of the fluorescent lamp are degraded, and the life span is short.

SUMMARY OF THE INVENTION

[0003] To solve the above problems, a feature of the present invention is to provide a white light source having an excellent electric field electron emission efficiency to thereby obtain a large emission current even at a low applied voltage, and having a very high density of electron emitters per unit area to thereby exhibit excellent luminous efficacy, and a fabrication method thereof.

[0004] In order to achieve the above feature, the present invention provides a white light source including a metal film used as a cathode, the metal film being formed on a lower substrate; a conductive polymer film pattern formed on the metal film; carbon nanotubes for emitting electrons, the carbon nanotubes being substantially vertically bound with the conductive polymer film pattern such that one end thereof is exposed above the surface of the conductive polymer film pattern; spacers installed on the metal film; and a transparent upper substrate on which a transparent electrode, to which a fluorescent body is attached, is formed, the transparent upper substrate being mounted on the spacers such that the fluorescent body faces the carbon nanotubes.

[0005] The fluorescent body is formed of $(3\text{Ca}_3(\text{PO}_4)_2\text{CaFCI/Sb,Mn})$, generating a white luminescence, or $\text{Y}_2\text{O}_3\text{:Eu}$, $\text{CeMaA}_{11}\text{O}_{19}\text{:Tb}$ and $\text{BaMg}_2\text{Al}_6\text{O}_7\text{:Eu}$, to generate a white luminescence based on three combined emission spectrums.

[0006] The present invention also provides a method of fabricating a white light source. A metal film used as a cathode is formed on a lower substrate. An insulation film pattern having a plurality of openings selectively exposing the metal film is formed on the metal film. The openings are filled with a conductive polymer film pat-

tern. Carbon nanotubes are scattered on the openings and sunken in the conductive polymer film pattern such that the carbon nanotubes vertically stand with one end exposed. The conductive polymer film pattern is hardened to bind the sunken carbon nanotubes with the conductive polymer film pattern. Spacers are installed on the insulation film pattern. A transparent upper substrate having a transparent electrode, to which a fluorescent body is attached, is mounted on the spacers such that the fluorescent body faces the carbon nanotubes, and the transparent upper substrate is sealed with the lower substrate.

[0007] The carbon nanotubes may be scattered on the openings by applying direct current bias or supersonic waves to the lower substrate. The fluorescent body is formed of $(3\text{Ca}_3(\text{PO}_4)_2\text{CaFCI/Sb,Mn})$ generating a white luminescence, or $\text{Y}_2\text{O}_3\text{:Eu}$, $\text{CeMaA}_{11}\text{O}_{19}\text{:Tb}$ and $\text{BaMg}_2\text{Al}_6\text{O}_7\text{:Eu}$ to generate a white luminescence based on three combined emission spectrums.

[0008] According to the present invention, a white light source having excellent luminous efficacy can be provided.

BRIEF DESCRIPTION OF THE DRAWINGS

[0009] The above feature and advantages of the present invention will become more apparent by describing in detail preferred embodiments thereof with reference to the attached drawings in which

FIG. 1 is a schematic sectional view for explaining a white light source according to an embodiment of the present invention; and

FIGS. 2 through 7 are schematic sectional views for explaining a method of fabricating a white light source according to the embodiment of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

[0010] Hereinafter, an embodiment of the present invention will be described in detail with reference to the attached drawings. The present invention is not restricted to the following embodiment, and many variations are possible within the spirit and scope of the present invention. The embodiment of the present invention is provided in order to more completely explain the present invention to anyone skilled in the art. In the drawings, the shapes of members are exaggerated for clarity and the same reference numerals denote the same members. Also, when a film is described as being on another film or a semiconductor substrate, it can be directly on the other layer or the semiconductor substrate or an interlayer film can exist therebetween.

[0011] The present invention provides a white light source using carbon nanotubes and a method of fabricating the same. It is known that a carbon nanotube is microscopically constituted such that a single carbon el-

ement is combined with three neighboring carbon atoms, a hexagonal ring is formed by the combination among the carbon atoms, and a plane composed of repeated hexagonal rings like a honeycomb is rolled to thereby form a cylindrical shape. The cylindrical structure is characterized by the diameter being usually several nanometers through several tens of nanometers, and the length being several tens through several thousands of times longer than the diameter.

[0012] Accordingly, the tip of a carbon nanotube has a diameter of several nanometers through several tens of nanometers, thereby realizing very high electric field electron emission efficiency. Therefore, a large amount of emission current can be obtained at a low applied voltage. In addition, carbon nanotubes can be aligned with a very high density per unit area so that a very high tip density can be achieved, thereby obtaining excellent luminous efficacy

[0013] FIG 1 is a schematic sectional view for explaining a white light source according to an embodiment of the present invention. Referring to FIG. 1, the white light source according to the embodiment of the present invention includes a metal film 200, used as a cathode, and a conductive polymer film pattern 400 on a lower substrate 100, and carbon nanotubes 500 vertically or obliquely planted in the conductive polymer film pattern 400.

[0014] The lower substrate 100 may be formed of silicon (Si), alumina (Al_2O_3), quartz or glass, but it is preferable that the lower substrate 100 is formed of glass which is suitable for a sealing process of completing a white light source. Since the metal film 200 serves to transmit current to the conductive polymer film pattern 400 bearing the carbon nanotubes 500, it is preferable that the metal film 200 is formed of a material of excellent conductivity. For example, the metal film 200 may be formed of a conductive material, for example, chrome (Cr), titanium (Ti), nitride titanium (TiN), tungsten (W) or aluminum (Al). However, since the conductive polymer film pattern 400 also exhibits conductivity, the metal film 200 can be omitted as necessary.

[0015] The conductive polymer film pattern 400 is sectioned in a predetermined pattern by an insulation film pattern 300. Accordingly, the carbon nanotubes 500 planted in the sectioned conductive polymer film pattern 400 are also grouped. Each group of carbon nanotubes 500 may constitute a single cell.

[0016] The carbon nanotubes 500 are planted in the conductive polymer film pattern 400 such that the tips of the carbon nanotubes 500 are exposed above the surface of the conductive polymer film pattern 400, and point toward the surface of a transparent electrode 800 used as an anode. An electric field applied to the transparent electrode 800 and the metal film 200 causes the electric field to concentrate at the tips of the carbon nanotubes 500, and thus electrons are emitted from the tips of the carbon nanotubes 500.

[0017] The emitted electrons collide with a fluorescent

body 900, which is attached to the transparent electrode 800 to face the carbon nanotubes 500, and thus the fluorescent body 900 radiates light. The fluorescent body 900 is patterned to have a predetermined shape. The transparent electrode 800 is formed of a transparent conductive material such as indium tin oxide (ITO) and is attached to a transparent substrate 700 formed of glass. The fluorescent body 900 may be formed of a fluorescent material, for example, $(3Ca_3(PO_4)_2CaFCI/Sb, Mn)$, generating a white luminescence, or a combination of fluorescent materials including, for example, $Y_2O_3:Eu$, $CeMg_{11}O_{19}:Tb$ and $BaMg_2Al_{16}O_{27}:Eu$, to generate a white luminescence based on the three combined emission spectrums.

[0018] The carbon nanotubes 500 are separated by a predetermined distance from the fluorescent body 900 by spacers 600 which are installed on the insulation film pattern 300. Each of the spacers 600 has a length of about 100-700 μm . The transparent substrate 700 is mounted on the spacers 600. For installation of the spacers 600, the fluorescent body 900 is patterned such that portions, at which the spacers 600 contact the transparent electrode 800, are exposed. The transparent substrate 700, to which the transparent electrode 800 is attached, is mounted on the spacers 600 and vacuum sealed with the lower substrate 100.

[0019] Since the diameter of the tip of each carbon nanotube 400 is very small - on average, several nanometers through several tens of nanometers, in an extreme case, as small as an atom, compared to the length of the carbon nanotube 400, electrons can be very effectively emitted from the tips in the presence of an applied electrical field. In other words, even if low voltage is applied between the metal film 200 or the conductive polymer film pattern 400 and the transparent electrode 800, a very high electric field can be formed at the tips of the carbon nanotubes 500 so that very effective emission of electrons can be achieved. In addition, since the carbon nanotubes 500 can be planted at very high density per unit area, the emission current caused by the electrons emitted from these carbon nanotubes 500 is very high.

[0020] The emitted electrons are transferred to the transparent electrode 800 by the electric field, which is applied between the transparent electrode 800, used as an anode, and the metal film 200 or the conductive polymer film pattern 400, used as a cathode. As the electrons collide with the fluorescent body 900, electrons in the fluorescent body 900 are excited to a high energy level and then de-excited to a lower energy level by emitting photons, thereby allowing the fluorescent body 900 to radiate light. The radiated light is emitted out through the transparent substrate 700. Here, as described above since the electron emission efficiency is high and the emission current due to the emitted electrons is also high, the current of electrons striking the fluorescent body 900 is very large. Accordingly, the luminance generated in the fluorescent body 900 is very high.

[0021] Although the white light source according to the embodiment of the present invention is substantially very simple and compact, it can emit white light with very high luminance, as described above. In addition, since it has a high electron emission efficiency, it can operate with very small voltage or very low current. Accordingly, this white light source can be used as a general illumination system, and if it is extremely miniaturized, it can be used as a portable one.

[0022] With reference to FIGS. 2 through 7, a method of fabricating the white light source according to the embodiment of the present invention as described above, will be described in detail.

[0023] FIG. 2 schematically shows the step of forming the conductive polymer film pattern 400 on the lower substrate 100. The metal film 200 used as a cathode is thinly formed on the lower substrate 100 having a large area for mass production. The lower substrate 100 may be formed of silicon, alumina, quartz or glass, but it is preferable that it is formed of glass to be suitable for sealing a white light source. The metal film 200 is formed by depositing a material of excellent conductivity, for example, chrome, titanium, nitride titanium, tungsten or aluminum, to a thickness of about 0.3-0.8 μm . The deposition is performed by a method of forming a thin film such as a thermal deposition method or a sputtering method.

[0024] An insulation film is deposited on the metal film 200 to a thickness of about 1.0-4.0 μm using an insulation material such as silicon oxide. The insulation film is deposited at a low temperature, for example, of below about 500°C, when the lower substrate 100 is formed of glass. This is for preventing deformation of the lower substrate 100 during the deposition of the insulation film.

[0025] Thereafter, the insulation film is patterned by photolithography, thereby forming the insulation film pattern 300 selectively exposing the underlying metal film 200. For example, photoresist is deposited to a thickness of about 1.5-2.0 μm and exposed and developed, thereby forming a photoresist pattern 350 selectively exposing the insulation film. Thereafter, the insulation film is selectively etched using the photoresist pattern 350 as an etching mask, thereby forming the insulation film pattern 300 selectively exposing the underlying metal film 200. The openings of the insulation film pattern 300 may be microscopic holes having a diameter of about 1-10 μm . The distance between the holes may be 3.0-15.0 μm . Then, the photoresist pattern 350 is removed by a stripping process.

[0026] FIG. 3 schematically shows the step of filling the insulation film pattern 300 with the conductive polymer film pattern 400. The microscopic holes of the insulation film pattern 300 are filled with liquid conductive polymer to contact the metal film 200 exposed by the insulation film pattern 300, thereby forming the conductive polymer film pattern 400. It is preferable that each of the openings is half filled with the conductive polymer. Accordingly, the conductive polymer film pattern 400

may be formed to a thickness of about 1-2 μm .

[0027] FIG. 4 schematically shows the step of planting the carbon nanotubes 500 in the conductive polymer film pattern 400. The carbon nanotubes 500 are scattered on and sunken in the conductive polymer film pattern 400. Here, since the carbon nanotubes 500 have a longish shape, most of them drop with one end directing downward. Accordingly, one end of the carbon nanotubes 500 first contacts the conductive polymer film pattern 400, and thus the carbon nanotubes 500 vertically sink into the conductive polymer film 400.

[0028] Since the microscopic openings filled with the conductive polymer film pattern 400 have a microscopic diameter of about 1-10 μm , the carbon nanotubes 500 obliquely or horizontally dropping are caught by the insulation film pattern 300. The carbon nanotubes 500 caught by the insulation film pattern 300 are slanted to one side and, substantially, vertically sunken in the conductive polymer film pattern 400.

[0029] To allow the carbon nanotubes 500 caught by the insulation film pattern 300 to be smoothly sunken in the conductive polymer film pattern 400, the lower substrate 100 may be shaken, or vibrated by supersonic waves. Alternatively, the carbon nanotubes 500 can be guided to be vertically sunken in the conductive polymer film pattern 400 by applying direct current (DC) bias to the lower substrate 100. By applying vibration or DC bias, the carbon nanotubes 500 caught by the insulation film pattern 300 can be vertically sunken in the conductive polymer film pattern 400. In other words the carbon nanotubes 500 are sunken in the conductive polymer film pattern 400 such that the tips thereof look directly or obliquely upward. The other ends of the carbon nanotubes 500 are exposed above the surface of the conductive polymer film pattern 400.

[0030] Meanwhile, it is preferable to scatter the carbon nanotubes 500 in a uniform density per unit area. For this purpose, a screen (not shown) is provided on the conductive polymer film pattern 400. The screen is shaken or vibrated after putting the carbon nanotubes 500 on the screen so that the carbon nanotubes 500 can be scattered through the holes of the screen. The holes of the screen are formed in a uniform density so that the carbon nanotubes 500 scattered through the holes can also be sunken in the conductive polymer film pattern 400 in a uniform density.

[0031] FIG. 5 schematically shows the step of hardening the conductive polymer film pattern 400 to bond the sunken carbon nanotubes 500. The conductive polymer film pattern 400 is hardened using the hardening characteristic of polymers constituting the conductive polymer film pattern 400. For example, when the conductive polymer film pattern 400 is composed of thermoset polymers, a hardening reaction is performed at a low temperature, for example, of below about 300°C to harden the conductive polymer film pattern 400. As a result, the sunken carbon nanotubes 500 are bound with the conductive polymer film pattern 400.

[0032] FIG. 6 schematically shows the step of installing the spacers 600 on the insulation film pattern 300. A plurality of spacers 600 of a length of about 100-700 μm are mounted on the insulation film pattern 300.

[0033] FIG. 7 schematically shows the step of forming the transparent electrode 800 and the fluorescent body 900 on the separated transparent substrate 700.

[0034] The transparent electrode 800 used as an anode is attached to the separated transparent substrate 700, for example, a glass substrate. The transparent electrode 800 is formed of a transparent conductive material such as ITO. Thereafter, the fluorescent body 900 is attached to the transparent electrode 800. The fluorescent body 900 may be formed of a fluorescent material, for example, $(3\text{Ca}_3(\text{PO}_4)_2\text{CaFCI/Sb,Mn})$, generating a white luminescence or fluorescent materials including, for example, $\text{Y}_2\text{O}_3\text{:Eu}$, $\text{CeMaA}_{11}\text{O}_{19}\text{:Tb}$ and $\text{BaMg}_2\text{Al}_6\text{O}_7\text{:Eu}$, to generate a white luminescence based on combined emission spectrums. The fluorescent body 900 may be patterned to allow the spacers 600 to contact the transparent electrode 800.

[0035] The separated substrate 700 having the fluorescent body 900 and the transparent electrode 800 is mounted on the spacers 600 such that the fluorescent body 900 and the transparent electrode 800 face the conductive polymer film pattern 400. By doing this, the tips of the carbon nanotubes 500, which are vertically bound with the conductive polymer film pattern 400, face the surface of the fluorescent body 900. After the transparent glass substrate 700 is mounted on the spacers 600, vacuum-sealing is performed.

[0036] For the carbon nanotubes 500 of the white light source fabricated through the above steps, the diameter of each tip is substantially very small at several nanometers through several tens of nanometers so that emission of electrons in an applied electric field can be achieved even with very low applied voltage.

[0037] As described above, the present invention can provide a white light source, which can obtain a large amount of emission current with low applied voltage, by using carbon nanotubes with ends having a very small diameter as electric field electron emission tips. In addition, the present invention can provide a white light source exhibiting excellent luminous efficacy since a very high density of tips per unit area can be achieved. Moreover, the processes of fabricating the white light source are simplified, thereby improving the yield and reliability of products. Accordingly, a next generation high efficient power saving white light source can be provided, replacing an existing fluorescent lamp or glow lamp. The white light source according to the present invention can be extremely miniaturized and has small power consumption so that it can be used as a portable white light source.

[0038] Although the invention has been described with reference to a particular embodiment, it will be apparent to one of ordinary skill in the art that modifications of the described embodiment may be made without de-

parting from the spirit and scope of the invention.

Claims

1. A white light source comprising:

a metal film used as a cathode, the metal film being formed on a lower substrate;
a conductive polymer film pattern formed on the metal film;
carbon nanotubes for emitting electrons, the carbon nanotubes being substantially vertically bound with the conductive polymer film pattern such that one end thereof is exposed above the surface of the conductive polymer film pattern;
spacers mounted on the metal film; and
a transparent upper substrate on which is formed a transparent electrode to which a fluorescent body is attached, the transparent upper substrate being mounted on the spacers such that the fluorescent body faces the carbon nanotubes.

2. A white light source as claimed in claim 1 wherein the lower substrate is formed of glass, quartz, alumina or silicon.

3. A white light source as claimed in claim 1 or 2 wherein the metal film is formed of chrome, titanium, titanium nitride, aluminium or tungsten.

4. A white light source as claimed in any preceding claim wherein the fluorescent body is formed of $(3\text{Ca}_3(\text{PO}_4)_2\text{CaFCI/Sb,Mn})$ generating a white luminescence, or $\text{Y}_2\text{O}_3\text{:Eu,CeMaA}_{11}\text{O}_{19}\text{:Tb}$ and $\text{BaMg}_2\text{Al}_6\text{O}_7\text{:Eu}$ to generate a white luminescence based on three combined emission spectrums.

5. A white light source comprising:

a metal film used as a cathode, the metal film being formed on a lower substrate;
an insulation film pattern provided on the metal film, the insulation film pattern having a plurality of openings selectively exposing the metal film;
a conductive polymer film pattern formed in the openings;
carbon nanotubes for emitting electrons, the carbon nanotubes being substantially vertically bound with the conductive polymer film pattern such that one end thereof is exposed above the surface of the conductive polymer film pattern;
spacers installed on the insulation film pattern; and
a transparent upper substrate on which is formed a transparent electrode to which a fluorescent body is attached, the transparent up-

per substrate being mounted on the spacers such that the fluorescent body faces the carbon nanotubes.

6. A white light source as claimed in claim 5, wherein the lower substrate is formed of glass, quartz, alumina or silicon.

7. A white light source as claimed in claim 5 or 6 wherein the metal film is formed of chrome, titanium, titanium nitride, aluminium or tungsten.

8. A white light source as claimed in any of claims 5 to 7, wherein the surface of the conductive polymer film pattern is lower than the surface of the insulation film pattern.

9. A white light source as claimed in any of claims 5 to 8, wherein the openings are holes having a diameter of about 1-10 μ m, and are separated from one another by a distance of about 3-15 μ m.

10. A white light source as claimed in any of claims 5 to 9, wherein the fluorescent body is formed of (3Ca₃(PO₄)₂CaFCI/Sb,Mn) generating a white luminescence, or Y₂O₃:Eu,CeMaA₁₁O₁₉:Tb and BaMg₂Al₁₆O₇:Eu to generate a white luminescence based on three combined emission spectrums.

11. A method of fabricating a white light source comprising the steps of:

forming a metal film used as a cathode on a lower substrate;

forming an insulation film pattern on the metal film, the insulation film pattern having a plurality of openings selectively exposing the metal film; filling the openings with a conductive polymer film pattern;

scattering carbon nanotubes on the openings and sinking the carbon nanotubes in the conductive polymer film pattern such that the carbon nanotubes vertically stand with one end being exposed;

hardening the conductive polymer film pattern to bind the sunken carbon nanotubes with the conductive polymer film pattern;

installing spacers on the insulation film pattern; and

mounting a transparent upper substrate on which is formed a transparent electrode to which a fluorescent body is attached on the spacers such that the fluorescent body faces the carbon nanotubes, and sealing the transparent upper substrate with the lower substrate.

12. A method as claimed in claim 11, wherein the lower

substrate is formed of glass, quartz, alumina or silicon.

13. A method as claimed in claim 11 or 12, wherein the metal film is formed of chrome, titanium, titanium nitride, aluminium or tungsten.

14. A method as claimed in claim 11, wherein the openings have a diameter of about 1-10 μ m and are separated from one another by a distance of about 3-15 μ m.

15. A method as claimed in any of claims 11 to 14, wherein the carbon nanotubes are scattered on the openings by applying direct current bias or super-sonic waves to the lower substrate.

16. A method as claimed in any of claims 11 to 15, wherein the openings are filled with the conductive polymer film pattern such that the surface of the conductive polymer film pattern is lower than the surface of the insulation film pattern.

17. A method as claimed in any of claims 11 to 16, wherein the fluorescent body is formed of (3Ca₃(PO₄)₂CaFCI/Sb,Mn) generating a white luminescence, or Y₂O₃:Eu,CeMaA₁₁O₁₉:Tb and BaMg₂Al₁₆O₇:Eu to generate a white luminescence based on three combined emission spectrums.

FIG. 1

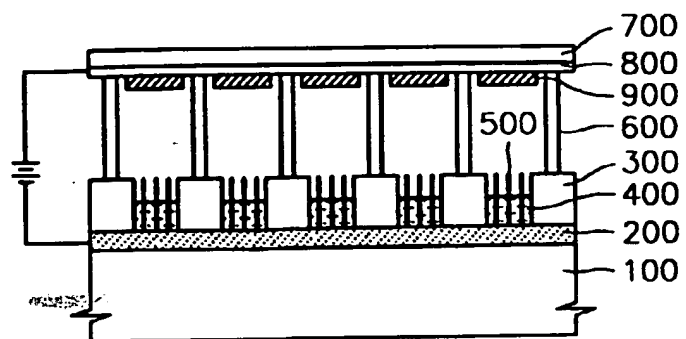


FIG. 2

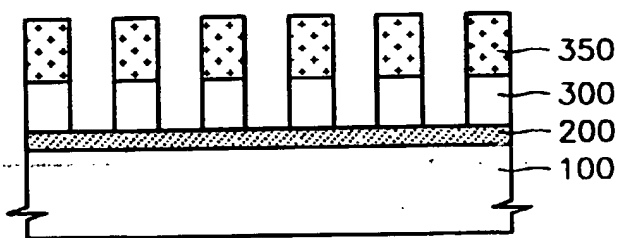


FIG. 3

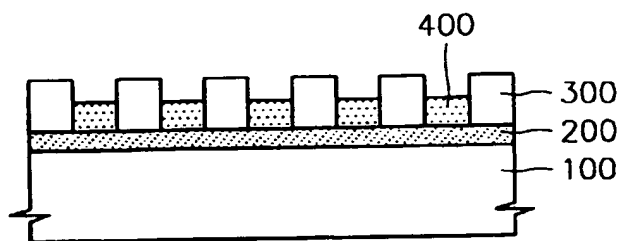


FIG. 4

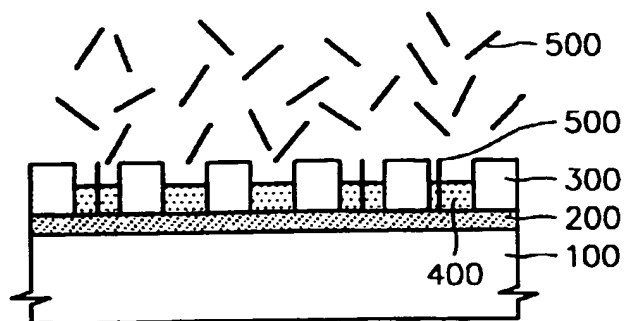


FIG. 5

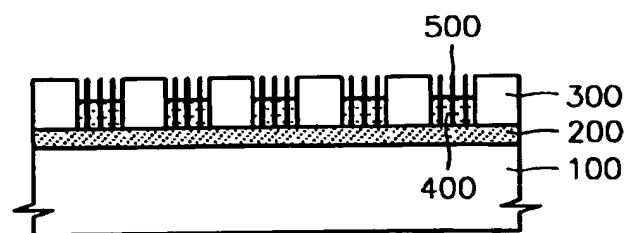


FIG. 6

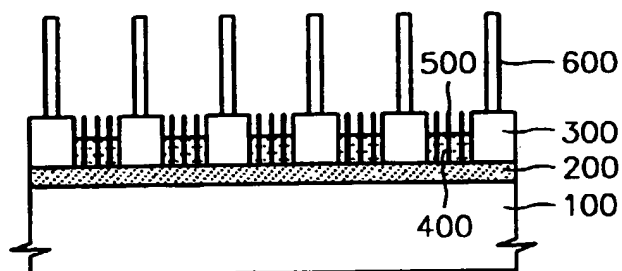


FIG. 7





European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 30 5140

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
P, X	EP 0 989 579 A (LUCENT TECHNOLOGIES INC ; UNIV NORTH CAROLINA (US)) 29 March 2000 (2000-03-29) * column 8, line 14 - column 10, line 23; claims 1-21 *	1, 3	H01J63/06 H01J9/24
Y		11	
Y	EP 0 913 508 A (CANON KK) 6 May 1999 (1999-05-06) * column 1, line 19-21; claims 22-25, 36 *	11	
A	WO 97 07531 A (DU PONT ; UNIV CALIFORNIA (US); SILZARS ARIS KENNETH (US); SPRINGER) 27 February 1997 (1997-02-27) * claims 1-13 *	1	
A	US 5 773 921 A (GROSSE-WILDE HUBERT ET AL) 30 June 1998 (1998-06-30) * column 6, line 52-56; claims 1-8 *	1	
A	WO 98 44526 A (ADVANCED TECH MATERIALS ; CANDESCENT TECH CORP (US)) 8 October 1998 (1998-10-08) * claims 1, 9, 22 *	1, 11	TECHNICAL FIELDS SEARCHED (Int.Cl.7) H01J
A	DE 196 02 595 A (BOSCH GMBH ROBERT) 31 July 1997 (1997-07-31)		
P, A	US 6 019 656 A (HA JEONG SOOK ET AL) 1 February 2000 (2000-02-01) * claim 1 *	11	
A	WO 94 28571 A (MICROELECTRONICS & COMPUTER) 8 December 1994 (1994-12-08) * claim 30 *	1	
A	EP 0 351 110 A (EMI PLC THORN) 17 January 1990 (1990-01-17) * claims 1-12 *	1	
		-/--	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 29 September 2000	Examiner Van den Bulcke, E
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons 8 : member of the same patent family, corresponding document	

EPO FORM 1503 03 82 (P4/C01)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 30 5140

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	WO 97 18577 A (DU PONT ; HERRON NORMAN (US); BLANCHET FINCHER GRACIELA BEAT (US);) 22 May 1997 (1997-05-22) * claims 1-14 *	11	
P,A	WO 99 66523 A (KITABATAKE MAKOTO ; SHIRATORI TETSUYA (JP); KUROKAWA HIDEO (JP); DE) 23 December 1999 (1999-12-23) *ABSTRACT*	1	
A	EP 0 905 737 A (ISE ELECTRONICS CORP) 31 March 1999 (1999-03-31) * column 8, line 33-47; claims 1-18 *	1,11	
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
Place of search THE HAGUE		Date of completion of the search 29 September 2000	Examiner Van den Bulcke, E
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

EPO FORM 1503 03/82 (P04/031)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 30 5140

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

29-09-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0989579 A	29-03-2000	JP 2000141056 A	23-05-2000
EP 0913508 A	06-05-1999	JP 11194134 A	21-07-1999
WO 9707531 A	27-02-1997	AU 696412 B	10-09-1998
		AU 7007596 A	12-03-1997
		CA 2229067 A	27-02-1997
		CN 1199503 A	18-11-1998
		DE 69605118 D	16-12-1999
		DE 69605118 T	25-05-2000
		EP 0845154 A	03-06-1998
		JP 11510951 T	21-09-1999
US 5773921 A	30-06-1998	DE 4405768 A	24-08-1995
		AT 186422 T	15-11-1999
		WO 9523424 A	31-08-1995
		DE 59507196 D	09-12-1999
		EP 0801805 A	22-10-1997
WO 9844526 A	08-10-1998	EP 0968509 A	05-01-2000
DE 19602595 A	31-07-1997	WO 9727607 A	31-07-1997
US 6019656 A	01-02-2000	NONE	
WO 9428571 A	08-12-1994	AU 5897594 A	20-12-1994
		CA 2164294 A	08-12-1994
		EP 0730780 A	11-09-1996
		JP 8510858 T	12-11-1996
		US 5600200 A	04-02-1997
		US 5686791 A	11-11-1997
		US 5675216 A	07-10-1997
		US 5659224 A	19-08-1997
		US 5703435 A	30-12-1997
EP 0351110 A	17-01-1990	AT 85729 T	15-02-1993
		CA 1305999 A	04-08-1992
		DE 68904831 D	25-03-1993
		DE 68904831 T	19-08-1993
		JP 2270247 A	05-11-1990
		JP 2806978 B	30-09-1998
		US 4969850 A	13-11-1990
WO 9718577 A	22-05-1997	AU 7678896 A	05-06-1997
		CN 1202271 A	16-12-1998
		DE 69604931 D	02-12-1999

EPO FORM P0469

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 30 5140

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

29-09-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9718577 A		DE 69604931 T	18-05-2000
		EP 0861499 A	02-09-1998
		JP 2000500905 T	25-01-2000
		US 5948465 A	07-09-1999
WO 9966523 A	23-12-1999	JP 2000090813 A	31-03-2000
EP 0905737 A	31-03-1999	JP 11111158 A	23-04-1999
		JP 2000036243 A	02-02-2000
		JP 11167886 A	22-06-1999

EPO FORM P/459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82